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18. (Amended) The method of claim 37, wherein the oligomers independently comprise subunits selected from the group consisting of deoxyribonucleotides, ribonucleotides, and analogs of deoxyribonucleotides or ribonucleotides; and any single oligomer comprises one or a combination of two or more of said different types of subunits.

*(3)* Claim 19 is amended as follows:

19. (Amended) The method of claim 37 wherein each of said oligomers forming said content addressable memory matrix  $T_{ij}$  comprises, in order from the 5' end to the 3' end, (a) an oligomer strand comprising a nucleotide sequence representing an  $i$ -th component of  $V$  selected from the group consisting of  $E_i$  and  $\underline{E}_i$  for  $i = 1$  to  $i = m$ , (b) an oligomer strand comprising a nucleotide sequence representing a  $j$ -th component of  $V$  selected from the group consisting of  $E_j$  and  $\underline{E}_j$  for  $j = 1$  to  $j = m$ , wherein  $j \neq i$ , and (c) a nucleotide sequence  $F$  that is not complementary to any sequence  $E_i$  or  $\underline{E}_i$  for  $i = 1$  to  $i = m$ .

(4) Claims 20-21 are canceled without prejudice or disclaimer.

*(5)* Claim 22 is amended as follows:

*(K 22)* 22. (Amended) The method of claim 37 wherein said single-stranded oligomers comprising a complete, stoichiometric set of  $E_i$  of step (c) and  $\underline{E}_i$  are anchored to a solid support.

(6) Claim 27 is amended as follows:

*Su B* 27. (Thrice Amended) The method of claim 9 wherein said operation of matrix or vector algebra is determining the inner product of two vectors  $V$  and  $W$ , and said method comprises:

*K3* (i) obtaining for each vector  $V$  and  $W$ , sets of single-stranded oligomers  $E_i$  and  $\underline{E}_i$  representing the components of the vector, wherein the concentrations of the oligomers  $E_i$  and  $\underline{E}_i$  are proportional to the absolute values of the amplitudes of the components they represent; and

also obtaining a set of single-stranded oligomers  $E_i$  and  $\underline{E}_i$  representing the components of vector  $W$  that are complementary to said oligomers representing vector  $W$ , wherein the relative concentrations of the oligomers representing  $W$  are proportional to the concentrations of their complementary oligomers in  $W$ ;

(ii) combining samples of the oligomers representing vector  $V$  with samples of the oligomers representing vectors  $W$  and  $\underline{W}$  in separate respective first and second reaction mixtures and measuring  $R_+$  and  $R_-$  rates of hybridization associated with the respective first and second mixtures, and obtaining a numerical value proportional to the inner product of the two vectors from a difference between said  $R_+$  and  $R_-$  rates of hybridization.

*K4* (7) Add new claim 36 as follows:

36. (New) The method of claim 9, wherein said operation of matrix or vector algebra includes obtaining an outer product matrix of two vectors  $V_i$  for  $i = 1, 2, \dots, m$ , and  $W_j$  for  $j = 1, 2, \dots, n$ , wherein said step of subjecting comprises obtaining a set of dimeric, single-stranded oligomers to represent an outer product of vectors  $V$  and  $W$ , each of said dimeric oligomers comprising (i) a first single-stranded oligomer sequence selected from the group consisting of  $E_i$  or  $\underline{E}_i$  for each  $i$ -th component of  $V$  for  $i = 1, 2, \dots, m$ , which oligomer is joined at its 3' end to the 5' end

of (ii) a second single-stranded oligomer sequence selected from the group consisting of  $E_j$  or  $\underline{E}_j$  for each  $j$ -th component of  $W$  for all  $j = 1, 2, \dots, n$ ,

wherein the step of detecting includes determining the concentration of said dimeric oligomers comprising oligomer sequences corresponding to the  $i$ -th component of  $V$  and the  $j$ -th component of  $W$ .

(8) Add new claim 37 as follows:

37. (New) A method for obtaining a data set  $V_i^b$  from an oligomer-based, content-addressable memory following input of a data set  $U_i^b$  that represents a portion of

$V_i^b$ ,

wherein data elements in the form of  $m$ -component vectors  $V = \sum_i V_i e_i$  are represented in the memory by a set of the oligomers  $E_i$  and  $\underline{E}_i$  that are a subset of all single-stranded oligomers and are in 1:1 correspondence with the basis vectors  $e_i$  for  $i = 1, 2, \dots, m$  in an abstract  $m$ -dimensional vector space;

wherein oligomers  $E_i$  and  $\underline{E}_i$  have complementary nucleotide sequences, with  $E_i$  oligomers representing the  $i$ -th component of  $V$  for which the amplitude  $V_i$  is positive, and  $\underline{E}_i$  representing the  $i$ -th component of  $V$  for which  $V_i$  is negative; and

wherein the concentration of each of oligomers  $E_i$  and  $\underline{E}_i$  is proportional to the absolute value of the amplitude  $V_i$  of the  $i$ -th component of  $V$ ;

the method comprising:

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(a) preparing a content-addressable memory representing memory matrix  $T_{ij}$  in which are stored data sets corresponding to vectors  $V_i^a$  for  $a = 1$  to  $a = n$ , where  $i = 1, 2, \dots, m$ , wherein  $T_{ij}$  is

the sum of all of the outer products  $V_i^a V_j^a$  for  $i \neq j$ , the preparing of the memory representing the matrix  $T_{ij}$ ;

comprising obtaining for each vector  $V^a$  a set of dimeric single-stranded oligomers, each of which comprises a first single-stranded oligomer sequence selected from the group consisting of  $E_i$  or  $\underline{E}_i$  for each  $i$ -th component of  $V^a$  for  $i = 1$  to  $i = m$ , and further comprises a second single-stranded oligomer sequence selected from the group consisting of  $E_j$  or  $\underline{E}_j$  for each  $j$ -th component of  $V^a$  for  $j = 1$  to  $j = m$ , except for  $i = j$ ; and then pooling said sets of dimeric oligomers obtained for each vector  $V^a$  for  $a = 1$  to  $a = n$  to form the set of oligomers of the content-addressable memory representing the matrix  $T_{ij}$ ;

(b) combining said pool of dimeric oligomers with a set of oligomers representing partial data Set  $U_i^b$  under conditions wherein oligomer sequences  $E_i^b$  and  $\underline{E}_i^b$  of data set  $U_i^b$  hybridize specifically to complementary sequences  $E_j$  and  $\underline{E}_j$  present in said memory pool oligomers; and obtaining an isolated set of monomeric oligomer strands  $X_i$  comprising the first single strand oligomer sequences  $E_i$  and  $\underline{E}_i$  of said memory pool of dimeric single stranded oligomers that hybridized specifically to said  $U_i^b$  oligomers, wherein said  $X_i$  oligomers do not further comprise said  $E_j$  and  $\underline{E}_j$  oligomers of the second single-stranded sequences of said memory pool oligomers that are complementary to said  $U_i^b$  oligomers;

(c) combining said set of  $X_i$  oligomers with a set of single-stranded saturating oligomers comprising a set of  $E_i$  and  $\underline{E}_i$  oligomers representing the complete set of basis vectors  $e_i$  for  $i = 1$  to  $m$ , wherein the  $E_i$  and  $\underline{E}_i$  oligomers are substoichiometric relative to said set of  $X_i$  oligomers, in that the number of oligomers in the set of  $X_i$  oligomers is greater than the number of saturating oligomers, so that complementary sequences hybridize to each other, denaturing the resulting duplex

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molecules, and isolating the subset of  $X_i$  oligomer that hybridized specifically to said  $E_i$  and  $\underline{E_i}$  sequences, to obtain a set of saturated  $X_i$  strands,  $S(X_i)$ ;

(d) repeating steps (b) and (c) iteratively, using the set of saturated  $X_i$  strands,  $S(X_i)$  obtained in each previous implementation of step (c) as the set of oligomers representing partial data set  $U_i^b$  employed in the subsequent implementation of step (b), until successive iterations yield the same set of oligomer strands  $X_i$  produced by step (b) that represents data set  $V_i^b$ .

*V. H. G.*  
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